

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
6 September 2002 (06.09.2002)

PCT

(10) International Publication Number  
WO 02/068382 A1

(51) International Patent Classification<sup>7</sup>: C07C 317/22

Kee-Young [KR/KR]; Inheon Apt. #101-602, 32-3,  
Bongcheon-Dong, Kwanak-Ku, Seoul 151-050 (KR).

(21) International Application Number: PCT/KR02/00199

(74) Agent: CHOI, Kyu-Pal; #401, Halla Classic Building  
4F, 824-11, Yeoksam-Dong, Kangnam-Ku, Seoul 135-080  
(KR).

(22) International Filing Date: 7 February 2002 (07.02.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:  
2001/9291 23 February 2001 (23.02.2001) KR

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,  
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,  
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,  
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK,  
LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,  
MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI,  
SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN,  
YU, ZA, ZM, ZW.

(71) Applicant (*for all designated States except US*): YON-  
SUNG FINE CHEMICAL CO. LTD. [KR/KR]; 51-301,  
Science Technology Park., Sungkyunkwan University,  
300, Chonchon-Dong, Changan-Gu, Suwon, Kyonggi-Do  
440-746 (KR).

(84) Designated States (*regional*): ARIPO patent (GH, GM,  
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),  
Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),  
European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR,  
GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent  
(BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR,  
NE, SN, TD, TG).

(72) Inventors; and

(75) Inventors/Applicants (*for US only*): HAM, Won-Hun  
[KR/KR]; Samik Apt. #12-1201, Cheongdam-Dong,  
Kangnam-Gu, Seoul 135-100 (KR). OH, Chang-Young  
[KR/KR]; 49-4, Seongsan-Dong, Mapo-Gu, Seoul  
121-250 (KR). KIM, Yong-Hyun [KR/KR]; 73-73,  
Hwagok 5-Dong, Kangseo-Gu, Seoul 157-015 (KR).  
LEE, Yiu-Suk [KR/KR]; Woosung Apt. #304-407, 105,  
Sadang-Dong, Dongjak-Gu, Seoul 156-090 (KR). LEE,

Published:

— with international search report

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

(54) Title: PROCESS FOR PREPARING SULFAMOYL-SUBSTITUTED PHENETHYLAMINE DERIVATIVES

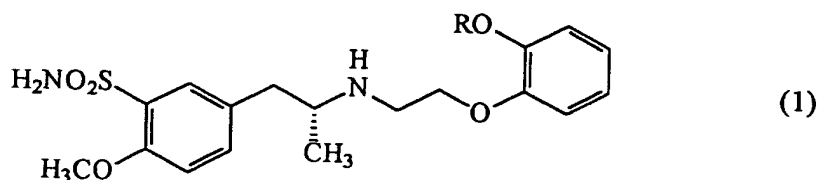
(57) Abstract: The present invention relates to a novel process for preparing sulfamoyl-substituted phenethylamine derivatives, specifically, 5-[2-[2-(2-alkoxy-phenoxy)ethylamino]-propyl]-2-methoxy-benzene sulfonamide of the formula (1).

WO 02/068382 A1

## PROCESS FOR PREPARING SULFAMOYL-SUBSTITUTED PHENETHYLAMINE DERIVATIVES

### TECHNICAL FIELD

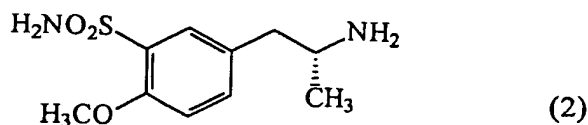
The present invention relates to a novel process for preparing sulfamoyl-substituted phenethylamine derivatives, specifically, 5-{2-[2-(2-alkoxyphenoxy)ethylamino]-propyl}-2-methoxy-benzene sulfonamide of the following formula:



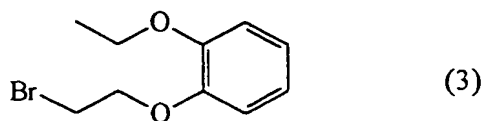
15 , wherein R represents C<sub>1-4</sub>alkyl, or hydrochlorides thereof.

### BACKGROUND ART

It is described in the United States Patent No. 5,447,958 that the compounds of the above formula (1) have excellent therapeutic effects against hypertension, congestive heart failure, angina pectoris or prostatic hypertrophy. In addition, in the above patent, a process for preparing the compounds of the formula (1) by reacting hydrochlorides of the compounds of the following formula:



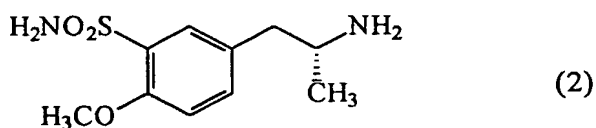
with the compounds of the following formula:



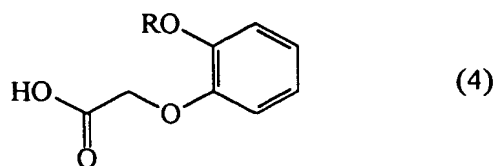
is disclosed. However, the above process has many drawbacks such as an extremely low yield (about 45%), a high production costs due to the purification by column chromatography and inapplicability in a large-scale production.

### DISCLOSURE OF THE INVENTION

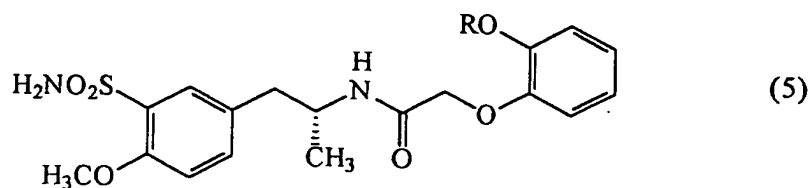
The present inventors have extensively studied to develop a novel preparation process in which the compounds of the formula (1) can be obtained in a high yield and can be easily purified. As a result, they found out that the preparation process of the compounds of the formula (1) by reacting compounds of the formula:



or hydrochlorides thereof with compounds of the formula:

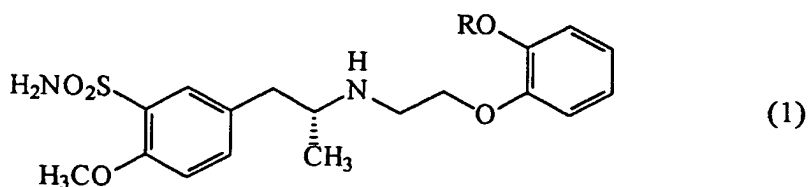


,wherein R represents C<sub>1-4</sub>alkyl, or acid chlorides or mixed anhydrides thereof, in the presence of a base, in a reaction solvent to obtain the compounds of the formula:



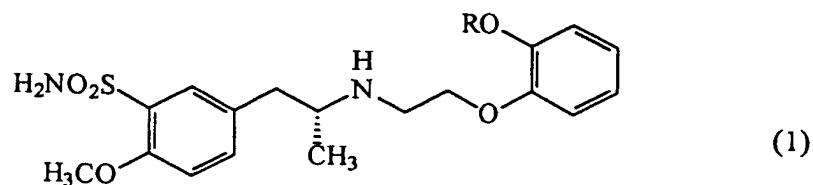
, wherein R is as defined above, and reducing the obtained compounds of the formula (5) with a reducing agent is not only economical but also efficient, because the compounds of the formula (1) can be obtained in a high yield, can be easily purified and can be prepared in a large scale, and completed the present invention.

Therefore, an object of the present invention is to provide a novel process for preparing sulfamoyl-substituted phenethylamine derivatives, specifically, compounds of the formula:



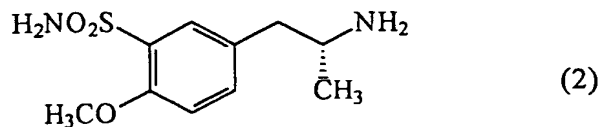
, wherein R represents C<sub>1-4</sub>alkyl, or hydrochlorides thereof.

The present invention relates to a process for preparing the compounds of the formula:

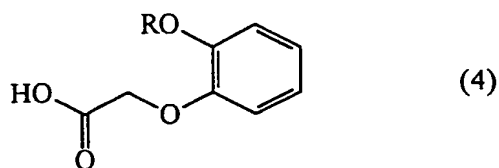


, wherein R represents C<sub>1-4</sub>alkyl, or hydrochlorides thereof, which comprises

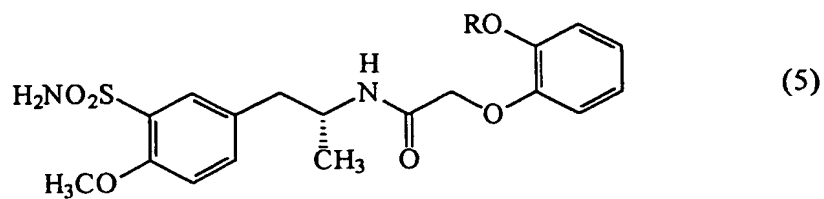
(i) reacting compounds of the formula:



or hydrochlorides thereof with compounds of the formula:



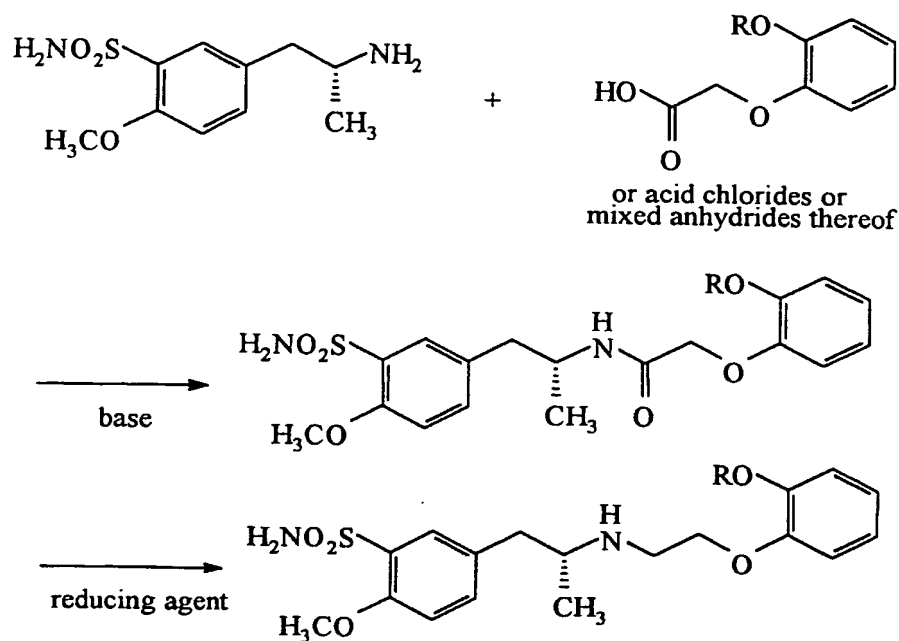
, wherein R is as defined above, or acid chlorides or mixed anhydrides thereof, in the presence of a base, in a reaction solvent to obtain compounds of the formula:



, wherein R is as defined above; and,

(ii) reducing the obtained compounds of the formula (5) with a reducing agent.

The above reaction may be depicted by the following reaction scheme:



Hereinafter, the present invention will be specifically explained. The process according to the present invention comprises the following two steps.

#### 5 **1st step: Preparation of compounds of the formula (5)**

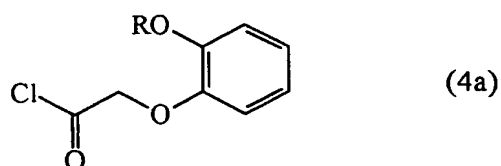
Compounds of the formula (5) are prepared by reacting compounds of the formula (2) or hydrochlorides thereof with compounds of the formula (4), or acid chlorides or mixed anhydrides thereof, in the presence of a base, in a reaction solvent. More specifically, the compounds of the formula (5) can be prepared by:

- 10 a) reacting compounds of the formula (2) or hydrochlorides thereof with acid chlorides of compounds of the formula (4);
- b) reacting compounds of the formula (2) or hydrochlorides thereof with mixed anhydrides of compounds of the formula (4); or,
- c) reacting compounds of the formula (2) or hydrochlorides thereof with
- 15 compounds of the formula (4).

In the above reaction, any conventional base such as trialkylamine, for example,

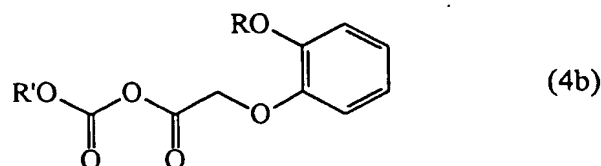
trimethylamine, triethylamine or diisopropylethylamine, or inorganic bases, for example,  $K_2CO_3$ ,  $Na_2CO_3$ ,  $KHCO_3$  or  $NaHCO_3$ , etc. may be employed and triethylamine is particularly preferable.

Acid chlorides of the compounds of the formula (4) employed in the process variant a) are represented by the following formula:

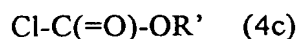


, wherein R represents  $C_{1-4}$ alkyl, which can be prepared by reacting the compounds of the formula (4) with an acid chloride, for example,  $SOCl_2$ ,  $PCl_3$ ,  $PCl_5$ ,  $POCl_3$  or oxalyl chloride, etc. A reaction solvent employable in the above process variant a) may be any polar or non-polar solvent which does not affect the reaction. Dimethylformamide (DMF), dimethylsulfoxide (DMSO), dimethylacetamide (DMA) or tetrahydrofuran (THF), etc. is preferable and THF is particularly preferable. A reaction temperature ranging from about 0 to 100 °C is preferable.

Mixed anhydrides of the compounds of the formula (4) employed in the process variant b) are represented by the following formula:



, wherein R represents  $C_{1-4}$ alkyl and R' represents alkyl, allyl or aryl, which can be prepared by reacting the compounds of the formula (4) with compounds of the formula:



, wherein R' is as defined above.

Preferable examples of the compounds of the formula (4c) include C<sub>1</sub>-alkylchloroformate such as methyl chloroformate, ethyl chloroformate or isobutyl chloroformate, phenyl chloroformate or allyl chloroformate, etc. A reaction solvent employable in the above process variant b) may be any polar or non-polar solvent which does not affect the reaction. DMF, DMSO, DMA or THF, etc. is preferable and THF is particularly preferable. A reaction temperature ranging from about -20 to 50 °C is preferable.

10 In the process variant c) wherein compounds of the formula (4) *per se* are employed, compounds of the formula (5) can be prepared by reacting compounds of the formula (2) or hydrochlorides thereof with compounds of the formula (4), in the presence of an acylating agent in addition to a base. Examples of the acylating agent employable in the above reaction include dicyclohexylcarbodiimide, diisopropylcarbodiimide, 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide, bis(2-oxo-3-oxazolidinyl)phosphinic chloride, benzotriazol-1-yloxy-tris(dimethylamino)phosphonium hexafluorophosphate, O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate, or O-7-azabenzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluoroborate, etc. Preferable is dicyclohexylcarbodiimide. A reaction solvent employable in the above process variant c) may be any polar or non-polar solvent which does not affect the reaction. DMF, DMSO, DMA or THF, etc. is preferable and DMF, DMSO or DMA is particularly preferable. A reaction temperature ranging from about 0 to 100 °C is preferable. Any additive, for example, dimethylaminopyridine, hydroxybenzotriazole or N-hydroxysuccinimide, etc. can be employed in the above process variant c). Such additive increases the reaction yield by suppression of side reactions.

In the first reaction step, an acylating agent, a base or an additive may be employed at a variable amount, preferably, at an amount of 1 to 3 equivalents, based on 1 equivalent of compounds of the formula (2) or hydrochlorides thereof. The reaction is performed while stirring for 1 to 24 hours. Then, the obtained compounds of the formula



(5) are filtered, and the filtrate is concentrated under reduced pressure, extracted with ethyl acetate, washed, dried, filtered and concentrated under reduced pressure and then, employed in the subsequent reaction step.

5 **2nd step: Preparation of compounds of the formula (1)**

Compounds of the formula (1) are prepared by reducing the compounds of the formula (5) obtained in the above first reaction step with a reducing agent in a reaction solvent. The reaction solvent employable in the above step may be any polar or non-polar solvent which does not affect the reaction. DMF, DMSO, DMA or THF, etc. is preferable and THF is particularly preferable. Examples of the reducing agent include lithium aluminum hydride, borane, diisobutylaluminum hydride, sodium borohydride-iodine or sodium borohydride-sulfate, etc. The reducing agent may be employed at an amount of about 2 to 6 equivalents. Preferably, a temperature of the reduction reaction ranges from about 40 to 80 °C and the reduction reaction is performed for 12 to 24 hours. Hydrochlorides of the compounds of the formula (1) can be obtained by addition of an anhydrous hydrochloric acid to the compounds of the formula (1).

The preparation process of the present invention is advantageous over the known process in that a higher yield can be obtained and the isolation step such as chromatography is not needed due to the easiness of purification.

**BEST MODE FOR CARRYING OUT THE INVENTION**

This invention will be better understood from the following examples. However, one skilled in the art will readily appreciate the specific materials and results described are merely illustrative of, and are not intended to, nor should be intended to, limit the invention as described more fully in the claims, which follow thereafter.

**Example 1: Synthesis of compounds of the formula (5) according to the process variant a)**

1.30 g (11 mmol) of thionyl chloride was added to 1.96 g (10 mmol, 1.0 eq.) of (2-ethoxy-phenoxy)acetic acid. The mixture was heated under reflux for 30 minutes and distilled under reduced pressure to obtain (2-ethoxy-phenoxy)acetyl chloride. 20 ml of THF and 2.78 ml (20 mmol, 2.0 eq.) of triethylamine were added to 2.44 g (10 mmol, 1.0 eq.) of (R)-(-)-5-[(2-amino-2-methyl)ethyl]-2-methoxybenzene sulfonamide and the reaction solution was cooled down to 0 °C. The obtained (2-ethoxy-phenoxy)acetyl chloride was added dropwise thereto and the whole mixture was stirred for 1 hour. The formed solids were filtered, and the filtrate was distilled under reduced pressure and extracted with 40 ml of ethyl acetate. Subsequently, the extract was washed with 1 N HCl, a saturated aqueous solution of sodium bicarbonate and brine, and dried over magnesium sulfate and then, filtered. The filtrate was distilled under reduced pressure to obtain 3.80 g (yield: 90%) of 2-(2-ethoxy-phenoxy)-N-[2-(4-methoxy-3-sulfamoyl-phenyl)-1-methyl-ethyl]-acetamide.

<sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz) 1.13 (d, 3H), 1.44 (t, 3H), 2.72 (dd, 1H), 2.85 (dd, 1H), 3.98 (s, 3H), 4.08 (m, 2H), 4.27 (m, 1H), 4.48 (dd, 2H), 5.06 (m, 3H), 6.87-7.02 (m, 5H), 7.34 (dd, 1H), 7.70 (d, 1H).

**Example 2: Synthesis of compounds of the formula (5) according to the process variant b)**

20 ml of THF and 1.10 ml (10 mmol, 1.0 eq.) of N-methylmorpholine were added to 1.96 g (10 mmol, 1.0 eq.) of (2-ethoxy-phenoxy)acetic acid. The reaction solution was cooled down to 20 °C and 1.29 ml (10 mmol, 1.0 eq.) of isobutyl chloroformate was added thereto and the whole mixture was stirred for 30 minutes. 2.44 g (10 mmol, 1.0 eq.) of (R)-(-)-5-[(2-amino-2-methyl)ethyl]-2-methoxybenzene sulfonamide in a mixture of 20 ml of THF and 1.10 ml (10 mmol, 1.0 eq.) of N-methylmorpholine was slowly added thereto and the whole mixture was stirred for 4 hours. The formed solids were filtered and the filtrate was distilled under reduced pressure and extracted with 40 ml of ethyl acetate. Subsequently, the extract was washed with 1 N HCl, a saturated aqueous solution of sodium bicarbonate and brine, dried over magnesium sulfate and filtered. The filtrate was distilled under reduced pressure to obtain 3.84 g (yield : 91%) of 2-(2-ethoxy-phenoxy)-N-

[2-(4-methoxy-3-sulfamoyl-phenyl)-1-methyl-ethyl]-acetamide.

NMR: as shown in the above example 1

**Example 3: Synthesis of compounds of the formula (5) according to the process variant c)**

50 ml of dimethylformamide was added to 2.44 g (10 mmol, 1.0 eq.) of (R)-(-)-5-[(2-amino-2-methyl)ethyl]-2-methoxybenzene sulfonamide and 1.96 g (10 mmol, 1.0 eq.) of (2-ethoxy-phenoxy)acetic acid and 1.35 g (10 mmol, 1.0 eq.) of hydroxybenzotriazole and 1.39 ml (10 mmol, 1.0 eq.) of triethylamine were sequentially added thereto and the reaction solution was cooled down to 0 °C. 2.06 g (10 mmol, 1.0 eq.) of dicyclohexylcarbodiimide was added thereto and the reaction solution was warmed up to room temperature and then, stirred for 24 hours. The formed solids were filtered and the filtrate was concentrated under reduced pressure and extracted with 50 ml of ethyl acetate. Subsequently, the extract was sequentially washed with water, 10% hydrochloric acid and a saturated aqueous solution of sodium bicarbonate, dried over magnesium sulfate and filtered. The filtrate was concentrated under reduced pressure to obtain 4.01 g (yield : 95%) of 2-(2-ethoxy-phenoxy)-N-[2-(4-methoxy-3-sulfamoyl-phenyl)-1-methyl-ethyl]-acetamide.

NMR: as shown in the above example 1

**Example 4: Synthesis of compounds of the formula (5) according to the process variant c)**

50 ml of dimethylformamide, 5.23 ml (30 mmol, 3.0 eq.) of N,N-diisopropylethylamine, 4.42 g (10 mmol, 1.0 eq.) of benzotriazol-1-yloxy-tris(dimethylamino)phosphonium hexafluorophosphate were sequentially added to 2.44 g (10 mmol, 1.0 eq.) of (R)-(-)-5-[(2-amino-2-methyl)ethyl]-2-methoxybenzene sulfonamide and 1.96 g (10 mmol, 1.0 eq.) of (2-ethoxy-phenoxy)acetic acid and the mixture was stirred for 12 hours. Upon the completion of the reaction, the formed solids were filtered, and the filtrate was distilled under reduced pressure and extracted with 40 ml of ethyl acetate. Subsequently, the extract was washed with 1 N HCl, a saturated aqueous solution of

sodium bicarbonate and brine, dried over magnesium sulfate and filtered. The filtrate was distilled under reduced pressure to obtain 4.05 g (yield : 96%) of 2-(2-ethoxy-phenoxy)-N-[2-(4-methoxy-3-sulfamoyl-phenyl)-1-methyl-ethyl]-acetamide.

NMR: as shown in the above example 1

5

#### Example 5: Synthesis of compounds of the formula (1)

50 ml of tetrahydrofuran was added to 4.22 g (10 mmol, 1.0 eq.) of 2-(2-ethoxy-phenoxy)-N-[2-(4-methoxy-3-sulfamoyl-phenyl)-1-methyl-ethyl]-acetamide. 379 mg (10 mmol, 4.0 eq.) of lithium aluminum hydride was added thereto and the whole mixture was stirred at 60 °C for 24 hours. Upon the completion of the reaction, 0.4 ml of water, 0.4 ml of 10 % NaOH aqueous solution and 1.2 ml of water were sequentially added to the reaction solution and then, filtered through celite. The filtrate was concentrated under reduced pressure to obtain 3.5 g of 5-{2-[2-(2-ethoxy-phenoxy)-ethylamino]-propyl}-2-methoxy-benzene sulfonamide. Thereto was added 10 ml of 2 M HCl solution in ethanol to obtain 3.56 g (yield : 80%) of 5-{2-[2-(2-ethoxy-phenoxy)-ethylamino]-propyl}-2-methoxy-benzene sulfonamide hydrochloride.

$[\alpha]_D^{20}$  4.0 (c=0.35, methanol)

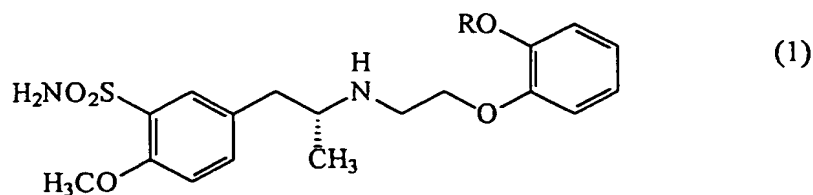
$^1\text{H}$  NMR (DMSO- $d_6$ , 500 MHz) 1.13 (d, 3H), 1.24 (t, 3H), 2.67 (dd, 1H), 3.27 (dd, 1H), 3.39 (m, 2H), 3.52 (m, 1H), 3.87 (s, 3H), 4.00 (q, 2H), 4.28 (t, 2H), 6.87-7.00 (m, 4H), 7.17 (s, 2H), 7.06-7.62 (m, 3H), 9.22 (s, 2H).

### INDUSTRIAL APPLICABILITY

According to the preparation process of the present invention, the sulfamoyl-substituted phenethylamine derivatives can be economically and efficiently prepared in a large scale, because a high yield can be obtained and purification can be performed without any isolation such as chromatography.

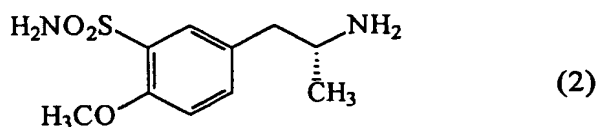
## WHAT IS CLAIMED IS:

1. A process for preparing the compounds of the formula:

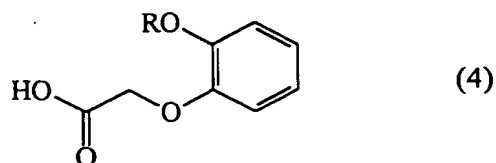


, wherein R represents C<sub>1-4</sub>alkyl, or hydrochlorides thereof, which comprises

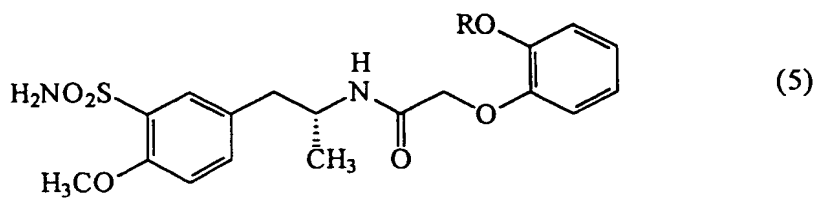
- (i) reacting compounds of the formula:



or hydrochlorides thereof with compounds of the formula:



, wherein R is as defined above, or acid chlorides or mixed anhydrides thereof, in the presence of a base, in a reaction solvent to obtain compounds of the formula:

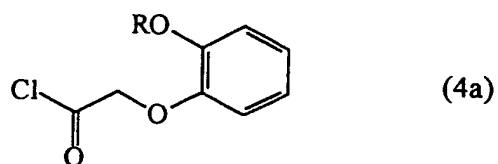


, wherein R is as defined above; and,

(ii) reducing the obtained compounds of the formula (5) with a reducing agent.

2. The process of claim 1, wherein the acid chlorides of the compounds of the formula (4) are represented by the formula:

5



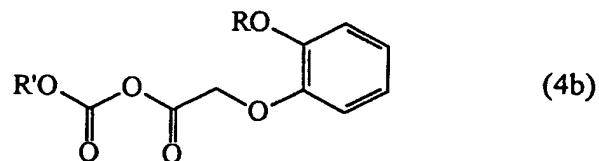
10

, wherein R represents C<sub>1-4</sub>alkyl.

3. The process of claim 2, wherein the compounds of the formula (4a) are prepared by reacting the compounds of the formula (4) with an acid chloride selected from the group consisting of SOCl<sub>2</sub>, PCl<sub>3</sub>, PCl<sub>5</sub>, POCl<sub>3</sub> and oxalyl chloride.

15

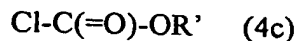
4. The process of claim 1, wherein the mixed anhydrides of the compounds of the formula (4) are represented by the formula:



, wherein R represents C<sub>1-4</sub>alkyl and R' represents alkyl, allyl or aryl.

25

5. The process of claim 4, wherein the compounds of the formula (4b) are prepared by reacting the compounds of the formula (4) with the compounds of the formula :



, wherein R' is alkyl, allyl or aryl.

30

6. The process of claim 5, wherein R' is methyl, isobutyl or phenyl.

7. The process of claim 1, wherein the compounds of the formula (2) or hydrochlorides thereof are reacted with the compounds of the formula (4), in the presence  
5 of an acylating agent in addition to the base to obtain the compounds of the formula (5) and the obtained compounds of the formula (5) are reduced with the reducing agent to obtain the compounds of the formula (1) or hydrochlorides thereof.

8. The process of claim 7, wherein the acylating agent is  
10 dicyclohexylcarbodiimide, diisopropylcarbodiimide, 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide, bis(2-oxo-3-oxazolidinyl)phosphinic chloride, benzotriazol-1-yloxy-tris(dimethylamino)phosphonium hexafluorophosphate, O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate or O-7-azabenzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluoroborate.

15

9. The process of any one of claims 1 to 8, wherein the base is trialkylamine or inorganic base.

10. The process of claim 9, wherein the trialkylamine is trimethylamine,  
20 triethylamine or diisopropylamine.

11. The process of any one of claims 1 to 8, wherein the reaction solvent is dimethylformamide (DMF), dimethylsulfoxide (DMSO), dimethylacetamide (DMA) or tetrahydrofuran (THF).

25

12. The process of any one of claims 1 to 8, wherein the reducing agent is lithium aluminum hydride, borane, diisobutylaluminum hydride, sodium borohydride-iodine or sodium borohydride-sulfate.



## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/KR02/00199

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC7 C07C 317/22

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**Minimum documentation searched (classification system followed by classification symbols)  
C07C7Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched  
KR : IPC as aboveElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
CAS On line**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	JP 2000-229901 A (Kato Yoshinori et al) 22.08.2000, see the whole document	1 ~ 12
Y	JP 62-114952 A (Fujikura Takashi et al) 26.05.1987, see the whole document	1 ~ 12
A	JP 02-306958 A (Ito Yasuo et al) 20.12.1990, see the whole document	1 ~ 12

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

## \* Special categories of cited documents:


- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

Date of the actual completion of the international search  
28 MAY 2002 (28.05.2002)

Date of mailing of the international search report  
29 MAY 2002 (29.05.2002)

## Name and mailing address of the ISA/KR

 Korean Intellectual Property Office  
920 Dunsan-dong, Seo-gu, Daejeon 302-701,  
Republic of Korea  
Facsimile No. 82-42-472-7140

## Authorized officer

KOH, Tae Wook  
Telephone No. 82-42-481-5545

